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European Patent Office
Office européen des brevets

⑪ Publication number:

0 279 493
A2

⑫

EUROPEAN PATENT APPLICATION

⑬ Application number: 88200261.1

⑮ Int. Cl.4: C07C 51/41, C07C 53/126,
C07C 65/05, C10M 129/28,
C10M 129/54

⑭ Date of filing: 11.02.88

⑯ Priority: 16.02.87 GB 8703549

⑰ Date of publication of application:
24.08.88 Bulletin 88/34

⑲ Designated Contracting States:
AT BE CH DE ES FR GB GR IT LI LU NL SE

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㉓ Process for the preparation of a basic salt, salt thus prepared and oil compositions containing such a salt.

㉔ Process for the preparation of basic alkaline earth metal salts of organic carboxylic acids, which comprises
(a) mixing the organic carboxylic acids with more than an equivalent amount of an alkaline earth metal hydroxide and/or oxide in a hydrocarbon solvent;
(b) introducing carbon dioxide into the mixture obtained in an amount of at least 0.5 equivalent carbon dioxide per equivalent of excess alkaline earth metal; and
(c) removing residual solids, if any, and an aqueous layer, if any, wherein the organic carboxylic acids comprise a blend of a C₈₋₃₀ alkyl salicylic acid and one or more alkanecarboxylic acid in which the alkyl moiety is branched and has from 4 to 40 carbon atoms.
Such a salt has dispersant properties and is suitable for use in lubricating oil and fuel compositions.

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PROCESS FOR THE PREPARATION OF A BASIC SALT, SALT THUS PREPARED AND OIL COMPOSITIONS
CONTAINING SUCH A SALT

The present invention relates to a process for the preparation of a basic alkaline earth metal salt of a blend of organic carboxylic acids, a salt thus prepared and to oil compositions containing such a salt.

The use of alkaline earth metal salts of organic carboxylic acids as additives for lubricating oil compositions is known. The said salts have a dispersant property so that, when applied in such 5 composition, they ensure that the inside of engine cylinders remains clean and that deposition of carbonaceous products on pistons and in piston grooves is counteracted, so that piston-ring sticking is prevented.

It is also known to prepare basic (or overbased) alkaline earth metal salts of such acids. The overbasing provides an alkaline reserve which, when applied in lubricating oil compositions, reacts with and neutralises 10 acidic compounds formed during the operation of the engine in which the composition is applied. Hence, sludge which may arise, is maintained dispersed due to the dispersant property of the salt while acids which would enhance sludge formation are neutralised.

In British patent specification No. 786,167, a process for the preparation of basic salts is described in which an organic acid is reacted with an excess of an alkaline earth metal oxide or hydroxide in an oil and 15 subsequently carbon dioxide is passed through the reaction mixture to yield basic salts. As suitable acids are mentioned substituted or unsubstituted aliphatic, cycloaliphatic and aromatic acids, comprising carboxylic acids, sulphur-containing acids, phosphoric acids, thio-acids, phenols and partial esters of sulphur-and phosphorus-containing acids.

In the technical field there is a desire to use products with a basicity as high as possible, i.e. the 20 relative proportion of the organic acid residue in the basic salt is as low as possible. The reason for this is that the costs of the product are mainly incurred by the costs of the organic acid.

The basicity of these products is generally expressed as a basicity index (BI), being defined as the 25 equivalents ratio of the total of alkaline earth metal to the total of organic acids. The prior art processes normally yield products having a BI of at most 10, and it was found that in the preparation of products having a BI that high or even higher using carboxylic acids a pronounced tendency to gelation occurs, thereby severely hindering the handleability of the products. Another problem was that when using certain acids only relatively low BI values were obtainable. It has now been found that when a blend of two specific types of carboxylic acids is used, a high BI value can be obtained without incurring gelation problems.

Accordingly, the present invention relates to a process for the preparation of a basic alkaline earth metal 30 salt of a blend of organic carboxylic acids, which comprises

(a) mixing the organic carboxylic acids with more than an equivalent amount of an alkaline earth metal hydroxide and/or oxide in a hydrocarbon solvent;
35 (b) introducing carbon dioxide into the mixture obtained in an amount of at least 0.5 equivalent carbon dioxide per equivalent of excess alkaline earth metal; and
(c) removing residual solids, if any, and an aqueous layer, if any, wherein the organic carboxylic acids comprise a blend of a C₈₋₃₀ alkyl salicylic acid and one or more alkanecarboxylic acids in which the alkyl moiety is branched and has from 4 to 40 carbon atoms.

Particularly preferred alkyl salicylic acids are those having at least 10 carbon atoms in the alkyl group, in particular from 12 to 26 carbon atoms.

40 The alkanecarboxylic acid used in the present invention is branched, and has a chain length long enough to avoid any solubility problems in oil compositions such as lubricants, whilst not being too long to incur problems in naphthenic or aromatic oil compositions. Preferably, the alkyl moiety has 8 to 28 carbon atoms, and in particular is a tertiary alkyl moiety. The tertiary carboxylic acids are especially suitable since when they are used very high BI values can be obtained. Very suitable acids include mixtures of tertiary 45 acids, such as those sold under the trade name "VERSATIC" acids. These acids are prepared by subjecting an olefin, e.g. a propylene oligomer, or a linear mixed olefin such as those sold under the trade name "SHOP" to a Koch reaction, yielding branched carboxylic acids. When mixtures of acids are used the preferred length of the alkyl moiety relates to the average length of the alkyl groups.

50 The ratios between the alkyl salicylic acid and the alkane carboxylic acid may vary within wide ranges. Advantageously the equivalent ratio of the alkyl salicylic acid to alkanecarboxylic acid ranges from 10:1 to 1:10, preferably from 4:1 to 1:4.

The alkaline earth metal salts prepared include magnesium, calcium, strontium and barium salts. Preferably, the alkaline earth metal applied is magnesium or calcium.

The reaction mixture prepared in step (a) of the present process suitably contains a promotor,

preferably an oxygen-containing organic solvent and optionally water. Suitable solvents include C₁₋₆alcohols, polyhydric alcohols such as glycol, propylene glycol, glycerol or 1,3-dihydroxypropane, ethers such as C₁₋₄ monoethers of glycol or propylene glycol, di isopropyl ether, 1,3- or 1,4-dioxane, or 1,3-dioxolane. Preferably the promotor is a C₁₋₆alcohol, in particular methanol. It will be appreciated that in industrial processes use 5 may be made of technical solvents, and that the use of technically pure promotors, such as methanol, might incur the presence of water. Hence, in such cases addition of water per se is not required since its addition is made implicitly by the addition of the promotor.

The preparation of the mixture according to step a) of the present process can be carried out in any convenient way, e.g. by mixing the alkaline earth metal hydroxide and/or oxide with the promotor, if used, 10 and adding the acids, whether or not in the presence of the promotor or a hydrocarbon solvent, to the resulting mixture. It is preferred to mix the blend of the acids and the alkaline earth metal hydroxide and/or oxide in a hydrocarbon solvent and subsequently add the promotor. The promotor may contain a substantial amount of water. Preferably the water content is adjusted such that the percentage of water in the mixture amounts to 0 to 10%w, based on the total liquids.

15 The hydrocarbon solvent can be selected from a wide variety of solvents. Suitable solvents include hydrocarbon oils, such as solvent-refined and/or hydrogenated lubricating oils having a kinematic viscosity of 3.5-35 mm²/s at 100°C. Preferably it is an aromatic hydrocarbon or a hydrocarbon fraction rich in aromatics, such as gasoline. Suitable hydrocarbon solvents are benzene, toluene, xylene or mixtures thereof, xylene being particularly preferred. The amount of the solvent is not critical. Promotor:solvent 20 volume ratios up to 1 can suitably be applied, preference being given to ratios in the range from 0.1 to 0.6.

The concentration of the organic carboxylic acid in the solvent or solvent mixture can vary within wide limits. Suitably the acids equivalent concentration of acids is from 0.01 to 1 molar equivalent/kilogram, preferably from 0.1 to 0.8, based on the combined weight of organic carboxylic acids and hydrocarbon solvent.

25 The amount of alkaline earth metal to be added in step a) should be at least 1 equivalent, and is preferably more, so that the subsequent carbon dioxide supply results in very high BI compounds. In that case, the amount of alkaline earth metal hydroxide and/or oxide added in step a) is preferably from 10 to 25 equivalents per equivalent acid.

30 The temperature at which step a) is carried out is not critical, and may be ambient temperature or elevated temperature. Suitable temperatures include 15-150°C.

In step b) the temperature is advantageously from 15 to 150°C, preferably from 30 to 75°C. In order to obtain the elevated temperature it may be necessary to employ elevated pressures, since the desired reaction temperature may be above the atmospheric reflux temperature of the reaction mixture. Suitable pressures include 1 to 15 bar abs. Higher pressures are possible, but merely add to the costs of the 35 process. The rate at which the carbon dioxide is introduced is advantageously from 0.05 to 1.0 equivalent carbon dioxide per equivalent acid per minute. The carbon dioxide introduction is conveniently carried out by passing carbon dioxide, or a mixture of carbon dioxide with an inert gas, such as air or nitrogen, through the reaction mixture under slightly higher pressure than the pressure prevailing in the reaction mixture. Higher pressures may be employed. Carbon dioxide will be absorbed in the reaction mixture and will react 40 with the alkaline earth metal compounds present therein forming a basic complex salt of the organic acid salt and carbonate, hydroxide and/or oxide. The amount of carbon dioxide to be taken up in step b) is to a certain extent dependent on the amount of alkaline earth metal added in step a) of the present process. Suitably the relative amount of carbon dioxide is somewhat less than the relative amount of alkaline earth metal hydroxide or oxide.

45 Preferably the introduction of carbon dioxide in step b) is stopped after 0.5 to 0.9 equivalent carbon dioxide per equivalent alkaline earth metal has been taken up. Conveniently, this corresponds with 5 to 23 equivalent carbon dioxide per equivalent acid.

It has been found that an ageing period between step b) and step c) can be advantageous, since it increases the BI of the resulting basic salt. Such an ageing period amounts suitably to at least 15 min. A 50 maximum period is generally imposed by practical and/or economical reasons, and is generally below 20 hours. Preferably the period between steps b) and c) is from 1 to 4 hrs.

The reaction mixture at the end of step b) may be worked up by any method known in the art. It may be subjected to a centrifuging treatment to remove solids comprising unreacted alkaline earth metal hydroxide and/or oxide and/or non-colloidal alkaline earth metal carbonate, if any. The resulting solution 55 may then be subjected to a liquid-phase separation. One liquid phase can be an aqueous layer which may contain the promotor when it is used, the other one is the hydrocarbon solvent plus the basic salts dispersed therein. It is also possible to reverse the above operations.

The present process can be used for the preparation of basic salts having a wide variety of basicity

indices. So, it would be possible to prepare basic salts having a relatively low BI e.g. from 1 to 10. The present process, however, is excellently suitable for preparing basic salts having a basicity index from 10 to 20.

5 The process described is a one-step process. However it is possible to integrate the process according to the present invention in a two-step process, in particular in a two-step process according to Applicants' UK patent application No. 8613815. Thereto, step (a) and (b) are carried out in two stages, a1, a2, b1 and b2 respectively, whereby the stages comprise:

(a1) preparing a mixture of one equivalent of the blend of the organic carboxylic acids and more than one equivalent of an alkaline earth metal hydroxide and/or oxide in a hydrocarbon solvent;

10 (b1) introducing carbon dioxide into the mixture obtained until at least 0.5 equivalent carbon dioxide per equivalent of excess alkaline earth metal has been taken up;

(a2) adding at least one further equivalent of alkaline earth metal hydroxide and/or oxide to the reaction mixture, so that the total amount of alkaline earth metal hydroxide and/or oxide is at least 10 equivalent;

15 (b2) resuming the introduction of carbon dioxide to the resulting mixture.

Between stages (b1) and (a2) and after (b2) ageing periods can be employed as is indicated in the above patent application. The ageing period after stage (b2) corresponds with the above-mentioned ageing period between steps (b) and (c).

20 The process according to the present invention can be carried out batchwise and also in a continuous manner.

The basic salts are excellent dispersant additives in oils. Therefore the present invention also provides oil compositions comprising a major amount of a base oil and a minor amount of a basic alkaline earth metal salt as described hereinbefore. Preferably the base oil is a lubricating base oil, and conveniently constitute more than 50%w of the composition. It can be selected from mineral lubricating oils of varying 25 viscosities, but it also includes a synthetic lubricant, such as ester-type lubricant or a polyolefin-type fluid, or a vegetable oil, or a grease.

Fuel compositions which are used in marine diesel engines usually contain some sulphur compounds. To neutralize the acidic compounds formed from these sulphur compounds a relatively high concentration 30 of the basic salt is employed. Preferably, these marine lubricating oil compositions contain from 5 to 30%w of basic alkaline earth metal salt. Lubricating oil compositions for road engines may contain lower concentrations. The amount of basic alkaline earth metal salt in these lubricating oil compositions is preferably from 0.01 to 5%w, in particular from 0.1 to 4.0%w.

Fuels, such as gasoline, kerosine, diesel fuel and gas oils, can also contain the above basic salts. The amount of these salts is similar to that in road engine lubricating oil compositions or lower; conveniently the 35 amount is from 0.001 to 5%w, in particular from 0.01 to 1.0%w.

The lubricating oil composition can be prepared by mixing a concentrate containing up to 60%w of a basic salt as described above in a lubricating oil, with a lubricating base oil to give the desired concentration. Such a concentrate is conveniently prepared by addition of a lubricating oil to the product obtained after completion of step c), and removal of any volatile hydrocarbon solvent, water and alcohol, if 40 present. The lubricating oil may be the same as the one indicated above as a suitable hydrocarbon solvent. The concentrate may conveniently contain a stabiliser, which is selected from a variety of organic compounds, such as those described in British patent specification No. 818,325. These compounds include mono- or polyhydric alcohols, alkyl amines and alkyl phenols.

The lubricating oil compositions may further contain a number of other additives, such as antioxidants, 45 foam inhibitors, corrosion inhibitors, viscosity index improvers, and pour point depressants, as can be established by a person skilled in the art. In particular, improved properties can be realised by addition of polyisobutene/succinic anhydride adducts, such as LUBAD 349 (sold by Lubrizol).

The invention will be illustrated by means of the following Examples.

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EXAMPLE 1

The following experiments were carried to show the synergistic effect of using a blend of organic carboxylic acids in the process of the present invention.

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Comparative Experiment A

1340ml of xylene was mixed with 640g of C₈-C₁₀ alkylsalicylic acid (ASA) (0.72 eq) and 400g (10.8 eq) calcium hydroxide. The Ca(OH)₂/ASA ratio was 15:1. After stirring for 1 hour at 40°C, 390 ml methanol was 5 added to the reaction mixture, and carbon dioxide was introduced at 50°C and a rate of 0.12 eq CO₂/eq acid.min. After an uptake of 9.3 eq CO₂/eq acid the reaction mixture gelated to become a solid mass. The BI of this solid mass could not be determined, but in theory could only have reached a value of around 10-11.

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Comparative Experiment B

This experiment used a mixture of tertiary carboxylic acids, sold under the trade name "VERSATIC 10", which consisted mainly of 3-methyl-octane-3-carboxylic acid together with minor amounts of 4-methyl-octane-4-carboxylic acid, 3-ethyl-heptane-carboxylic acid and 4-ethyl-heptane-4-carboxylic acid. The acid 15 number of this acids mixture was 5.8 meq/g. One equivalent of this mixture was added to xylene, yielding an acid concentration of 0.2 meq/g. Calcium hydroxide was added in an amount of 20 eq. After stirring for 1 hour, methanol was added to the reaction mixture at 44-55°C to give 20% vol based on total liquid. CO₂ was introduced at a rate of 0.40 eq CO₂/eq acid.min, until an uptake of 12 eq CO₂/eq acid. After settling of the 20 two liquid phases now present, the methanol-water layer was removed: From the xylene layer the residual solids were removed by centrifugation. The BI of the calcium salt present in the xylene layer was 5.3.

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The procedure of comparative experiment B was repeated with a mixture of the ASA of experiment A and the V10 acids of experiment B. The relative amounts of the various reactants were 1 equivalent ASA per 1 equivalent V10 acid. The concentration of the acids in xylene was 0.4 meq H⁺/g, and the relative amount of calcium hydroxide was 10 equivalent. Methanol, containing 3% vol water, was added in a 30 quantity of 14.4% vol, based on the xylene. Carbon dioxide was introduced into the reaction mixture at a rate of 0.12 eq(eq acid.min at a temperature of 50-55 °C. After an CO₂ uptake of 7.1 eq(eq acid, the reaction mixture was worked up as described in experiment B, yielding a product having a BI of 8.8, without gelling.

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Experiment 2

A two-step process was carried out by following the procedure of Experiment 1 with the following amounts of reactants: ASA to V10 acids equivalents ratio is 7.3; the equivalent ratio of calcium hydroxide to 40 acid is 10:1; the acid concentration, the composition of the solvents mixture, temperature and the CO₂ introduction rate were as described in Experiment 1. After a CO₂ uptake of 7.88 eq(eq acid, the reaction was interrupted for 15 minutes, calcium hydroxide was subsequently added in an amount of 7 eq(eq acid, the reaction mixture was stirred for one hour and the CO₂ introduction was resumed at 50°C. After a total uptake of 12.1 eq CO₂ (in both steps) the reaction mixture was treated as described in Experiment B, 45 yielding a product with a BI of 14.8.

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The procedure of Experiment 2 was repeated. After stopping of the CO₂ introduction at the second stage, i.e. after an uptake of 12.1 eq CO₂, the reaction mixture was subjected to an ageing period of 16 hours while stirring at 50°C continued. After the work-up procedure as described in Experiment B a product with a BI of 15.1 was obtained.

From these experiments it is apparent that whereas the use of ASA or V10 acids alone yields 55 unsatisfactory results, the use of a combination of these acids yield good products with a high BI and without incurring gelation problems. The use of an ageing period increases the BI of the product obtained.

EXAMPLE 2

In this Example a combination of ASA and tertiary carboxylic acids mainly having 19 carbon atoms (V19 acids) was used. The carboxylic acids were prepared by subjecting C₈ alpha-olefins to a Koch reaction.

5 The product of this reaction is a mixture of highly branched acids based on the C₈ olefin and its dimer which is formed during the reaction. The acid number of the mixture was 2.28 meq/g. Experiment 4 was carried out as a one step process as Experiment 1. Experiment 5 was carried out as Experiment 4 but with the use of a 16hr ageing period as described in Experiment 3. Experiments 6 and 7 were carried out as Experiment 3, i.e. with the use of a 16 hours ageing period.

10 The acids ratio, the CO₂-uptake in the various steps and the BI of the products obtained are indicated in Table 1 below.

TABLE 1

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| Exp. No. | ASA/V19 (eq/eq) | CO ₂ uptake (eqCO ₂ /eq acid) | | | BI product (eq/eq) |
|-------------|--------------------|---|--------|-------|--------------------------|
| | | 1 step | 2 step | total | |
| 4 | 1:1 | - | - | 12.5 | 13.7 |
| 5 | 1:1 | - | - | 12.5 | 14.2 |
| 6 | 1:1 | 7.9 | 4.1 | 12.0 | 13.0 |
| 7 | 7:3 | 7.9 | 4.6 | 12.5 | 14.6 |

The product of Experiments 4-7 were introduced into a mineral lubricating oil, the mixtures obtained were subjected to vacuum distillation to remove xylene to yield concentrates. The calcium contents and the kinematic viscosities of the concentrates at 100°C are indicated below.

TABLE 2

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| Prod. of Exp. No. | Ca content %w | V _k 100 (mm ² /s) |
|----------------------|------------------|--|
| 4 | 10.1 | 15.1 |
| 5 | 10.1 | 12.7 |
| 6 | 9.7 | 10.2 |
| 7 | 9.8 | 14.3 |

EXAMPLE 3

The procedure of Experiment 2 was followed in the following experiments 8 and 9. The differences between the latter experiments and Experiment 2 are as follows. The acids used in an equivalent ratio 1:1 were ASA and a mixture of tertiary carboxylic acids obtained as those described in Example 2 and having an acid number of 1.83 meq/g. In both steps an amount of calcium hydroxide of 12 eq/eq acid was added, i.e. the total calcium hydroxide amounted to 24 eq. In the work-up procedure residual solids were removed by filtration over a filter aid and not by centrifugation. Reaction conditions were as in Experiment 2. The CO₂ uptake and the BI's of the resulting products are given in Table 3.

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TABLE 3

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| Exp. No. | CO ₂ uptake (eqCO ₂ /eq acid) | | | BI product (eq/eq) |
|----------|---|--------|-------|--------------------|
| | 1 step | 2 step | total | |
| 8 | 9.4 | 7.7 | 17.1 | 18.31 |
| 9 | 9.4 | 9.6 | 19.0 | 18.95 |

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The products of these experiments were taken up in a lubricating oil as described in Example 2. The calcium contents and the kinematic viscosity are indicated below.

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TABLE 4

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| Prod. of Exp. No. | Ca content (%w) | V _k 100 (mm ² /s) |
|----------------------|--------------------|--|
| 8 | 9.9 | 11.6 |
| 9 | 9.9 | 12.9 |

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Comparative Experiment C

To show that linear carboxylic acids are not suitable for the present process, a combination of stearic acid and ASA was used in a 1:1 equivalent ratio in the following experiment.

To 96.1 g of ASA and 32.0 g of stearic acid in 435.3 g of xylene 141.9 g of calcium hydroxide (17 eq/eq acid) was added together with 91.7 g of methanol and 2.8 g of water. This reaction mixture was subjected to a reaction as described under comparative Experiment B, and at a CO₂ uptake of 8.2 eq/eq acid the CO₂ introduction was interrupted due to severe gelation of the reaction mixture.

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Claims

1. Process for the preparation of basic alkaline earth metal salts of organic carboxylic acids, which comprises
 - 5 (a) mixing the organic carboxylic acids with more than an equivalent amount of an alkaline earth metal hydroxide and/or oxide in a hydrocarbon solvent;
 - (b) introducing carbon dioxide into the mixture obtained in an amount of at least 0.5 equivalent carbon dioxide per equivalent of excess alkaline earth metal; and
 - (c) removing residual solids, if any, and an aqueous layer, if any;
- 10 characterised in that the organic carboxylic acids comprise a blend of a C₈₋₃₀ alkyl salicylic acid with one or more alkanecarboxylic acid in which the alkyl moiety is branched and has from 4 to 40 carbon atoms.
- 15 2. Process according to claim 1, in which the alkyl salicylic acid has 12 to 26 carbon atoms in the alkyl group.
- 15 3. Process according to claim 1 or 2, in which the alkyl moiety of the alkanecarboxylic acid is a tertiary alkyl moiety containing 8 to 28 carbon atoms.
- 15 4. Process according to any one of claims 1-3, in which the equivalent ratio of the C₈₋₃₀alkylsalicylic acid:alkane carboxylic acid ranges from 10:1 to 1:10.
- 20 5. Process according to any one of claims 1-4, in which the alkaline earth metal is calcium or magnesium.
- 20 6. Process according to any one of claims 1-5, in which the amount of alkaline earth metal hydroxide and/or oxide added in step (a) is from 10 to 25 equivalents per equivalent acid.
- 25 7. Process according to any one of claims 1-6, in which the mixture in step (a) further comprises an oxygen-containing organic solvent, and optionally water.
- 25 8. Process according to claim 7, in which the oxygen-containing organic solvent is a C₁₋₆alcohol.
- 25 9. Process according to any one of claims 1-8, in which the mixture of the blend of the organic carboxylic acids and alkaline earth metal hydroxide and/or oxide is prepared by mixing the acids and the alkaline earth metal hydroxide and/or oxide in the hydrocarbon solvent
- 30 10. Process according to claim 9, in which the hydrocarbon solvent is selected from benzene, toluene, xylene, or a mixture thereof.
- 30 11. Process according to any one of claims 1-10, in which the introduction of carbon dioxide in step (b) is carried out at a rate of 0.05 to 1.0 equivalent carbon dioxide per equivalent acid per minute.
- 35 12. Process according to any one of claims 1-11, in which an ageing period of from 0.25 to 20 hours is present between steps (b) and (c).
- 35 13. Process according to any one of claims 1-12, in which steps (a) and (b) are carried out in two stages, the stages comprising:-
 - (a1) preparing a mixture of the blend of organic carboxylic acids with more than an equivalent amount of an alkaline earth metal hydroxide and/or oxide in a hydrocarbon solvent;
 - (b1) introducing carbon dioxide into the mixture obtained until at least 0.5 equivalent carbon dioxide per equivalent of excess alkaline earth metal has been taken up;
 - (a2) adding at least one further equivalent of alkaline earth metal hydroxide and/or oxide to the reaction mixture, so that the total amount of alkaline earth metal hydroxide and/or oxide is at least 10 equivalent;
 - (b2) resuming the introduction of carbon dioxide to the resulting mixture.
- 45 14. Basic alkaline earth metal salt of a blend of organic carboxylic acids whenever prepared according to any one of claims 1-13.
- 45 15. Oil composition comprising a major amount of a base oil and a minor amount of a basic alkaline earth metal salt as claimed in claim 14.

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